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(54) Title: CANISTERS FOR USE IN METERED DOSE INHALERS

(57) Abstract: A canister suitable for use in metered dose inhalers and fitted with a metering valve characterised in that its walls are formed of a laminate comprising a first layer which is composed of a metal and a second layer which is composed of a strengthening material.



**WO 02/072449 A1**

### Canisters for use in Metered Dose Inhalers

This invention relates to an improved canister for use in metered dose inhalers (MDI's), especially to a canister which may contain and dispense pressurised formulations of pharmaceutical substances in propellants, for example, hydrofluoroalkane propellants.

Metered dose inhalers are a widely used system for delivery of pharmaceutical substances to the lung and upper airways. As such they have become established as a delivery system for pharmaceuticals for the treatment of respiratory disorders such as asthma and chronic obstructive pulmonary disease (COPD).

Metered dose inhalers typically comprise a canister fitted with a valve such as a metering valve which is fitted into a suitable channelling device to permit inhalation. Suitable channelling devices comprise, for example a valve actuator and a cylindrical or cone-like passage through which medicament may be delivered from the filled canister via the metering valve to the nose or mouth of a patient e.g. a mouthpiece actuator. In a typical arrangement the valve stem is seated in a nozzle block which has an orifice leading to an expansion chamber; the expansion chamber has an exit orifice which extends into the mouthpiece.

Metered dose inhalers operate by dispensing a metered volume of formulation by means of the metering valve from a bulk supply of formulation contained within the canister which generally consists of a suspension or solution of a substance in a liquefied propellant gas. In the past the preferred propellant gases were CFC's such as P11, P12 and 114. However, following the discovery that CFC's are capable of causing depletion of atmospheric ozone alternative propellants have been developed. The currently most favoured alternative propellants are hydrofluoroalkanes, especially 1,1,1,2-tetrafluoroethane (HFA134a) and 1,1,1,2,3,3,3-heptafluoro-n-propane (HFA227).

The medical community has been engaged for many years now in the process of formulating old and new medicines in the alternative propellants. It is now generally recognised that because of certain chemical and physical differences between the old and new propellants certain other modifications to the MDI system have been considered advisable. For example, the surfactants previously commonly used as dispersing agents in suspension based CFC formulations are not effective since they are inadequately soluble in hydrofluoroalkanes without the addition of a co-solvent to

the formulation. Efforts are being made to develop novel surfactants which are soluble in hydrofluoroalkanes (see e.g. WO96/09816 of Glaxo Inc). A system has also been developed in which the surfactant is omitted altogether (see e.g. WO93/11743 of Glaxo Group).

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Certain suspension formulations of pharmaceutical products in hydrofluoroalkanes have demonstrated a tendency to suffer drug losses due to deposition of drug substances on the canister walls. Also the possibility of interaction between the formulation and the canister walls cannot always be excluded. For this reason canister coatings have been developed e.g. coatings of a fluorocarbon polymer optionally in combination with a non-fluorocarbon polymer (see e.g. WO96/32151 of Glaxo Wellcome Inc). Coatings of epoxy-phenol resins are described in WO00/30608 (Chiesi). WO00/78286 (3M) describes coatings including a thin film of glass. Processes employed in order to apply and cure such coatings which involve use of elevated temperature can cause distortion of conventional aluminium canisters. Accordingly aluminium canisters have been developed with certain strengthening features such as thicker walls and ellipsoidal base, however these are relatively expensive to manufacture.

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Stainless steel is known as an alternative strong material for manufacture of canisters however lengths have to be taken to avoid corrosion e.g. as described in WO00/73170 (Boehringer Ingelheim) and use of special alloys is also expensive.

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We have now invented a canister suitable for use in metered dose inhalers which eliminates or substantially mitigates a number of the disadvantages of prior art canisters.

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Thus according to the invention we provide a canister suitable for use in metered dose inhalers and fitted with a metering valve characterised in that its walls are formed of a laminate comprising a first layer which is composed of a metal and a second layer which is composed of a strengthening material.

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Such canisters are advantageous in that the laminate is capable of providing equivalent or greater strength than thicker walled aluminium cans at a more reasonable cost. Furthermore, problems associated with possible interaction between the metal and the contents of the can or the atmosphere may be avoided.

Laminate when used in this specification will be understood to mean a material comprising two or more layers each of which make a contribution to the structural integrity of the canister when attached together.

- 5 Metals suitable for use according to the invention include pure metal or metal alloys which have optionally been pre-treated or processed e.g. galvanised, annealed, plated or coated to improve their properties. Preferably the metal is selected from the list consisting of aluminium, steel, copper, brass, tin and chromium, especially aluminium or steel, particularly aluminium.

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The strengthening material will be a material that when used in conjunction with the metal layer forms a product of greater strength than a similar conventional product not prepared from a laminated material. The strengthening material may have an inherent ability to resist stress or the ability may be realised when used in conjunction with the metal layer.

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One or more of the following materials may be used as a strengthening layer:

- i) a metal or an alloy e.g. aluminium, steel, iron, copper, tin, brass or chromium; or
- 20 ii) a metal oxide e.g. chromium oxide (which has good abrasive resistance and additionally is resistant to corrosion), alumina or tantalum oxide; or
- iii) a plastics material such as: polyester, epoxy resin, phenolic resin, phenoxy resin, epoxy-phenol resin, polypropylene, acrylonitrile-styrene copolymer, acrylonitrile-styrene-butadiene copolymer, high-impact polystyrene, nylon, polyacetal, polycarbonate, polytetrafluoroethylene, polyethylene terephthalate, an unsaturated
- 25 nitril resin, polyvinylchlorides, polyurethanes or polyphenylene oxides; optionally reinforced with a suitable filler material e.g. carbon fibres, filaments of glass, metal, boron, aluminium silicate, calcium carbonate, talc or barium sulphate; or
- iv) a ceramic e.g. silicon oxide; or
- v) a carbide (e.g. a carbide of a metal such as iron, calcium, tungsten or silicon or
- 30 boron).

The strengthening layer may be made from the same material as the metal (e.g. the laminate may be formed from two or more aluminium layers), although preferably the strengthening layer is usually selected from a different material to that of the first

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The laminate may optionally include one or more other layers such as:

- i) a bonding layer e.g. a layer of adhesive such as an epoxy resin, thermoset adhesive or cyanoacrylate;
- ii) a layer e.g. of a metal oxide which may be used to effect or improve adhesion between other layers such as between the metal layer and the strengthening layer;
- 5   iii) a coating layer e.g. applied to part or all of the internal and/or external surface of the laminate so as to serve one or more of the following purposes:
  - a) reduction of interaction of a layer of laminate with the atmosphere (e.g. in the case of the coating layer applied to the external surface);
  - b) reduction of interaction of a layer of laminate with the contents of the  
10   canister (e.g. in the case of the coating layer applied to the internal surface);
  - c) reduction of deposition of particulate matter from the contents of the canister onto a layer of laminate (e.g. in the case of the coating layer applied to the internal surface).

15   The above optional layers may perform more than one function, for example, adhesives may also distribute stress at the bond point and/or resist moisture and or corrosion. Similarly the oxide layer may be resistant to corrosion. Alternatively, a coating layer applied to an internal surface may reduce the interaction between the layer of laminate and the contents of the canister and also reduce the deposition of  
20   particulate matter from the contents of the canister onto the internal surface of the laminate.

A first particular embodiment is provided wherein the second layer of strengthening material is a plastics material such as polyester, polypropylene, acetonitrile-styrene  
25   copolymer, high-impact styrene, nylon or polyacetal, optionally reinforced with a material selected from carbon fibres, filaments of glass, metal, boron, aluminium silicate, calcium carbonate, talc and barium sulphate, and wherein the first metal layer is composed of a material selected from aluminium and steel and characterised in that the first metal layer forms the internal surface of the canister and is optionally  
30   coated with one or more coating layers. An adhesive layer may optionally be used between the plastics material layer and the first metal layer.

Preferably the plastics material is reinforced. More preferably the plastics material is reinforced with a material selected from carbon fibres and metal.

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A second particular embodiment is provided wherein the second layer of strengthening material is a plastics material such as polyester, polypropylene, acetonitrile-styrene copolymer, high-impact styrene, nylon or polyacetal, optionally

reinforced with a material selected from carbon fibres, filaments of glass, metal, boron, aluminium silicate, calcium carbonate, talc and barium sulphate, and wherein the second layer is optionally coated with one or more coating layers and the first metal layer is composed of a material selected from aluminium and steel and  
5 characterised in that the first metal layer forms the external surface of the canister and is optionally coated with one or more coating layers. An adhesive layer may optionally be used between the plastics material layer and the first metal layer.

Preferably the plastics material is reinforced. More preferably the plastics material is  
10 reinforced with a material selected from carbon fibres and metal.

According to a third particular embodiment of the invention we provide a canister, suitable for use in metered dose inhalers, characterised in that its walls are formed of a laminate wherein the laminate comprises a first metal layer of material forming the  
15 external surface of the canister selected from the list of metals consisting of: aluminium, brass, copper, chromium, iron, tin and steel, optionally coated with one or more coating layers, and a second layer of material selected from the list of strengthening materials consisting of aluminium, brass, copper, chromium, chromium oxide, iron, tin and steel, optionally coated with one or more coating layers.

20 The invention also provides a canister suitable for use in metered dose inhalers characterised in that its walls are formed of a laminate wherein the laminate comprises a first metal layer of material forming the internal surface of the canister selected from the list of metals consisting of: aluminium, brass, copper, chromium,  
25 iron, tin and steel, optionally coated with one or more coating layer, and a second layer of material selected from the list of strengthening materials consisting of aluminium, brass, copper, chromium, chromium oxide, iron, tin and steel, optionally coated with one or more coating layers.

30 According to the third and fourth particular embodiments the laminate may comprise further layers e.g. where the second layer is chromium a layer of chromium oxide may be present between the first and second layer and optionally an adhesive layer may be present.

35 Preferably the first metal layer is composed of stainless steel, aluminium or an alloy thereof, most preferably aluminium or stainless steel, especially aluminium.

Preferably the second layer of strengthening material is steel.

Thus in one embodiment of the invention the laminate is a laminate comprising one layer of steel and one layer of aluminium, optionally coated with one or more coating layers.

- 5 In a particularly preferred embodiment of the invention the wall of the canister is formed from a laminate comprising a layer of steel sandwiched between two layers of aluminium (i.e. is a laminate of one layer of steel and two layers of aluminium). Optionally the internal and/or external aluminium walls are provided with one or more coating layers. Preferably the internal wall of the canister will be coated with e.g. a  
10 fluorocarbon polymer coat such as PTFE or FEP; or a mixture of a fluorocarbon polymer with a non-fluorocarbon polymer such as PTFE or FEP and polyethersulphone; or an epoxy-phenol resin.

- The steel for use according to the invention will generally consist of a ferrous alloy  
15 with greater than 50% iron and one or more of the following: 0.1 to 10 wt% carbon (C), 0 to 10 wt% manganese (Mn), 0 to 10 wt% copper (Cu), 0 to 10 wt% silicon (Si), 0 to 50 wt% nickel (Ni), 0 to 10 wt% vanadium (V), 0 to 10 wt% niobium (Nb), 0 to 10 wt% aluminium (Al), 0 to 10 wt% molybdenum (Mo), 0 to 50 wt% chromium (Cr), 0 to 5 wt% phosphorus (P), 0 to 5 wt% sulphur (S), 0 to 30 wt% tungsten (W), 0 to 20  
20 wt% cobalt (Co), 0 to 10 wt% columbium (Cb), 0 to 10wt% oxygen (O), 0 to 10 wt% nitrogen (N) which may desirably be employed to improve the properties of the steel.

Stainless steel is a ferrous alloy as defined above which contains at least 12% chromium.

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Where the steel is sandwiched between two layers of metals, preferably the steel used will be mild steel which is a ferrous alloy as defined above containing up to 2% carbon.

- 30 The total thickness of the laminate for the finished product is preferably in the range 0.2-2.5mm, especially 0.2-2.0mm. When the laminate comprises a first layer which is metal and a strengthening layer which is metal the thickness of each layer is independently in the range 0.1-1.0mm. When the laminate is a laminate of one layer of steel and one layer of aluminium or stainless steel the thickness of the layer is  
35 preferably in the range 0.1-1.0mm. When the laminate is a laminate of one layer of steel and two layers of aluminium (or one layer of aluminium and one layer of stainless steel) the thickness of the sheets is preferably in the range 0.1-1.0mm. When the laminate comprises a strengthening layer selected from a plastics material

preferably the thickness of the said layer will be in the range 0.2-2.0mm especially, 0.8-1.5mm. The thickness of the laminate, or the constituent sheets thereof, may be reduced slightly with respect to the starting material when it is drawn into a canister.

- 5 As mentioned above, laminates may be prepared using an adhesive e.g. an epoxy resin with a molecular weight in the range 5000 to 30,000 (for further details see EP 0612 608). The adhesive will usually require curing e.g. by heating. Pressure sensitive adhesives (PSA) may also be used as an agent to bond the component layer together. The PSA may be activated by heating after assembly of the laminate
- 10 and processing using pressure lamination rollers (for further details please see USP 3970 496). This method is especially suitable when the laminate is composed of at least two layers which are a metal e.g. wherein the lamina comprises a layer of aluminium or stainless steel and a layer of steel or a sandwich as described above.
- 15 Alternative methods of preparing laminates include: surface activation of some or all the component lamina e.g. by using materials which when subjected to a mechanical force, such as pressure rollers with raised portions, some molecules/atoms of the two materials migrate and bind the said materials together, alternatively the migration of some molecules/atoms of the two material may be promoted by heating; electro
- 20 deposition e.g. electroplating with tin (for future details please see USP 5 298 149); or vapour deposition.

Canisters may be prepared from laminates by hot and/or cold drawing, extrusion or moulding. For further information on these processes please see Materials Science

25 and Engineering - an introduction by William D. Callister, Jr. (2nd edition), published by Wiley.

As mentioned above, canisters may be coated on their internal surface with a polymer e.g. a fluorocarbon polymer optionally in combination with a non-

30 fluorocarbon polymer as described in WO 96/32151.

Suitable fluoropolymers include polytetrafluoroethylene (PTFE), ethylenetetrafluoroethylene (ETFE), polyvinylidene fluoride (PVDF), perfluoroalkoxyalkane (PFA), polyvinyl fluoride (PVF), polychlorotrifluoroethylene

35 (PCTFE) and fluorinated ethylenepropylene (FEP). Fluorocarbon polymers are marketed under trademarks such as Teflon®, Tefzel®, Halar® and Hostaflon®, Polyflon® and Neoflon®. Grades of polymer include FEP DuPont 856-200, PFA



DuPont 857-200, PTFE-PES DuPont 3200-100, PTFE-FEP-polyamideimide DuPont 856P23485, FEP powder DuPont 532, and PFA Hoechst 6900n.

5 Fluorinated polymers which have a relatively high ratio of fluorine to carbon such as perfluorocarbon polymers e.g. PTFE, PFA and FEP are preferred.

Suitable copolymers comprise from 1 to 99%, preferably from 5 to 95% by weight of fluorinated polymer. Suitable copolymers include copolymers of tetrafluoroethylene (TFE) with PFA, TFE with hexafluoropropylene (HFP) (available as FEP 6107 and  
10 FEP 100 from DYNEON), VDF with HFP (commercially available as Viton A), TFE with perfluoro(propyl vinyl ether) (available as PFA 6515N from DYNEON), a blend of TFE, hexafluoropropylene and vinylidene fluoride (available commercially as THV 200G from DYNEON), HOSTAFORM X329<sup>TM</sup> (Hoechst) which is a 5% PTFE/Acetal blend, HOSTAFORM C9021TF which is a 20% PTFE/Acetal blend, PTFE/PBT  
15 blends (for example, LNP WL4040), and PTFE/PBT/silicone blends (for example, LNP WL4540).

The fluorinated polymer may be blended with non-fluorinated polymers such as polyamides, polyimides, polyamideimides, polyethersulphones, polyphenylene  
20 sulphides and amine-formaldehyde thermosetting resins. These added polymers improve adhesion to the canister walls, especially in the case of canisters lined with aluminium. Preferred polymer blends include PTFE/FEP/polyamideimide, PTFE/polyethersulphone and FEP/benzoguanamine.

25 The preferred polymer is a blend of polyethersulphone (PES) and polytetrafluoroethylene (PTFE). Another preferred polymer for coating is pure FEP (fluorinated ethylene propylene). Another preferred polymer for coating is pure PFA.

Canisters may be coated by the means known in the art for metal coating. For  
30 example, a metal such as aluminium or stainless steel may be pre-coated as a coil and cured before being stamped or drawn out into the can shape. Alternatively, pre-formed canisters may be sprayed inside with formulations of the coating polymer and then cured. The preformed canisters may also be dipped in the polymer coating formulation and cured, thus becoming coated on the inside and out. The polymer  
35 may also be formed in situ at the can walls using plasma polymerisation of the fluorocarbon monomers. The appropriate curing temperature is dependent on the polymer chosen for the coating and the coating method employed. However for coil coating and spray coating temperatures in excess of the melting point of the polymer

are typically required, for example, about 50°C above the melting point, for up to 20 minutes. For the named preferred and particularly preferred polymers, curing temperatures in the range of around 300-400 °C e.g. around 350-380 °C are suitable.

- 5 Alternative coatings include: epoxy resin, phenolic resin, phenoxy resin, epoxy-phenol resin, epoxy-phenol-novolac obtained by glycidylation of phenol-formaldehyde (novolac) condensation products, and epoxy-cresol-novolac obtained by glycidylation of o-cresol formaldehyde (o-cresol novolac) condensation products see WO00/30608.

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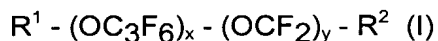
Other suitable coatings comprise linear non-cross-linked fluorinated polymers. In one aspect, the coating compound comprises a functional grouping which is capable of anchoring the compound to the surface thereof. As a first example, the compound may be an organo-phosphate such as a phosphate based perfluoroether derivative.

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Typically, the compound is a phosphoric ester.

In one first such embodiment, the interfacial surface has a compound disposed thereon having the general formula (I):

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wherein  $R^1$  comprises a fluoro-alkyl functional group;

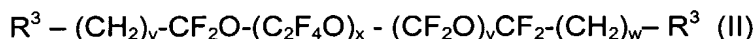
x and y are such that the molecular weight of the compound is in the range 350-1000; and

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$R^2$  comprises a phosphoric ester functional group.

In a second such embodiment, the interfacial surface has a compound disposed thereon having the general formula (II):

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wherein  $R^3$  comprises  $-(OCH_2-CH_2)_z-OPO(OH)_2$ ;

x, y and z are such that the molecular weight of the compound is in the range 900-2100; and v and w independently represent 1 or 2.

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In one preferred embodiment, v and w are both 1. In a second preferred embodiment v and w are both 2.

Alternatively in a second embodiment the compounds may be an organo-silane derivative such as a silane derivative of perfluoropolyoxyalkane, e.g. a silane derivative of perfluoropolyoxyalkane having a molecular weight in the range 1600-1750. Examples include perfluoropolyoxyalkanes having functional groups of the type  $-\text{CONR}^4\text{R}^5$  wherein  $\text{R}^4$  and  $\text{R}^5$  may be independently selected from hydrogen, or a silyl ether (e.g.  $\text{SiR}_t(\text{OR})_{3-t}$ ) wherein  $\text{R}$ =hydrogen or  $\text{C}_{1-8}$ alkyl and  $t=0$  to 2) as described in USP 4 746 550 which is incorporated herein by reference.

The synthesis of compounds of formula (I) and (II) may readily be determined by reference to EP687533 which describes similar compounds. EP338531 also provides information on the preparation of compounds of this type. Methods of preparing organo-silane polymeric compounds of the type described above may readily be determined by reference to USP 4 746 550.

Whilst not wishing to be bound by any theory, it is believed that the phosphate or silane moiety of the compounds of formula as described above reacts with the surface of the component to anchor the compound to the surface. Thus, when in use, the per-fluorinated end of the compound is presented to the pharmaceutical formulation and so provides a highly fluorinated surface.

Other suitable coatings include siloxanes such as dimethyl siloxane which in one aspect, may be applied by plasma polymerisation processes.

One suitable means of applying a fluorine-containing coating is by plasma coating, for example, by a  $\text{CF}_4$  or fluorine ion plasma coating technique. The plasma coating may consist of a fluorinated polymer laid down on the surface of the component by polymerisation or by modification of a hydrocarbon-containing pre-coating on the surface by interchange of hydrogen ions in the material with fluorine ions. The coating process typically takes place in a vacuum at ambient temperature. The components to be coated are placed inside a chamber which is evacuated. The fluorine monomer or fluorine source is introduced into the chamber at a controlled rate. The plasma is ignited within the chamber and maintained for a given time at a chosen power setting. For plasma polymerization typically temperatures in the range of about  $20^\circ\text{C}$  to about  $100^\circ\text{C}$  may be employed. At the end of the treatment the plasma is extinguished, the chamber flushed and the products retrieved. In the polymerisation process, a thin layer of plasma polymer will be bonded to the surface.

In one embodiment the internal wall of the canister may comprise a coating layer of high-nitril resin which is a co-polymer comprising an unsaturated nitril compound e.g. acrylonitrile or methacrylonitrile and an unsaturated copolymer e.g. an unsaturated aromatic compound, a diene compound an unsaturated ether compound or an unsaturated ester such as styrene,  $\alpha$ -methylstyrene, butadiene, isoprene, methylacrylate, ethylacrylate, methylmethacrylate and ethylmethacrylate, wherein said co-polymer contains greater than 50 percent by weight of the unsaturated nitril unit. The nitril resin layer may be manufactured by methods such as multilayer blow moulding, multilayer injection-blow moulding and multilayer injection moulding. Optionally an adhesive layer may be used between the nitril resin layer and the metal layer. The nitril resin layer has the advantage of being resistant to chemical attack and strong.

An alternative coating is a thin layer of glass which may be deposited by gas vapour deposition on the internal wall of the canister.

The coating thickness will typically be in the range 0.1 micron to 1 mm, e.g. 1-100 microns especially 1 to 25 microns.

Coatings may be applied in one or more coats.

Use of coatings, for example, plastics coatings may not be necessary if the surface, especially the internal surfaces of the canister are non-metallic, for example, a plastics material.

In use, the canister will be fitted with a valve, e.g. a metering valve, usually located in a cap covering the mouth of the canister. The metering valve is designed to deliver a metered amount of the formulation per actuation and incorporate a gasket to prevent leakage of propellant through the valve. Suitable valves are commercially available from manufacturers well known in the aerosol industry, for example, from Valois, France (eg. DF10, DF30, DF60), Bepak plc, UK (e.g. BK300, BK356, BK357) and 3M-Neotechnic Ltd, UK (eg. Spraymiser™). The DF31 valve of Valois, France is also suitable.

Valve metering volumes of 25  $\mu$ l, 50  $\mu$ l, 63  $\mu$ l or 100  $\mu$ l are typical.

Other valve systems include, but are not limited to, wedge gate valve systems, double-disc gate valve systems, globe and angle valve systems, swing check valve systems, end cock valve systems, and other like valve systems.

- 5 Valve materials, especially the metering chamber, will preferably be manufactured of a material which is inert to and resists distortion by the contents of the formulation, especially when the contents include ethanol. Particularly suitable materials for use in manufacture of the metering chamber include polyesters e.g. polybutyleneterephthalate (PBT) and acetals, especially PBT.

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Materials for use in the manufacture of the metering chamber and/or the valve stem may desirably be fluorinated, partially fluorinated or impregnated with fluorine containing substances in order to resist drug deposition.

- 15 Valve seals, especially the gasket seal, and also the seals around the metering chamber, will preferably be manufactured of a material which is inert to and resists extraction into the contents of the formulation, especially when the contents include ethanol.

- 20 Gaskets may comprise any suitable elastomeric material such as, for example, low density polyethylene, chlorobutyl, black and white butadiene-acrylonitrile rubbers, butyl rubber, neoprene, EPDM (a polymer of ethylenepropylenediene monomer) (eg as described in WO95/02651) and TPE (thermoplastic elastomer; eg as described in WO92/11190).

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The elastomeric material may either comprise a thermoplastic elastomer (TPE) or a thermoset elastomer which may optionally be cross-linked.

- 30 Typically, the sealing ring, which seals the valve to the can, and/or second sealing ring comprise an elastomeric material. The ring is typically resiliently deformable.

The sealing ring may be formed by cutting a ring from a sheet of suitable material. Alternatively, the sealing ring may be formed by a moulding process such as an injection moulding, a compression moulding or a transfer moulding process.

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The sealing ring may also comprise a thermoplastic elastomer blend or alloy in which an elastomeric material is dispersed in a thermoplastic matrix. The elastomers may optionally additionally contain conventional polymer additives such as processing

aids, colorants, tackifiers, lubricants, silica, talc, or processing oils such as mineral oil in suitable amounts.

5 Suitable thermoset rubbers include butyl rubbers, chloro-butyl rubbers, bromo-butyl rubbers, nitrile rubbers, silicone rubbers, fluorosilicone rubbers, fluorocarbon rubbers, polysulphide rubbers, polypropylene oxide rubbers, isoprene rubbers, isoprene-isobutene rubbers, isobutylene rubbers or neoprene (polychloroprene) rubbers.

10 Suitable thermoplastic elastomers comprise a copolymer of about 80 to about 95 mole percent ethylene and a total of about 5 to about 20 mole percent of one or more co-monomers selected from the group consisting of 1-butene, 1-hexene, and 1-octene as known in the art. Two or more such copolymers may be blended together to form a thermoplastic polymer blend.

15 Another suitable class of thermoplastic elastomers are the styrene-ethylene/butylene-styrene block copolymers. These copolymers may additionally comprise a polyolefin (e.g. polypropylene) and a siloxane.

20 Thermoplastic elastomeric material may also be selected from one or more of the following: polyester rubbers, polyurethane rubbers, ethylene vinyl acetate rubber, styrene butadiene rubber, copolyether ester TPE, olefinic TPE, polyester amide TPE and polyether amide TPE.

25 Other suitable elastomers include ethylene propylene diene rubber (EPDM). The EPDM may be present on its own or present as part of a thermoplastic elastomer blend or alloy, e.g. in the form of particles substantially uniformly dispersed in a continuous thermoplastic matrix (e.g. polypropylene or polyethylene). Commercially available thermoplastic elastomer blend and alloys include the SANTOPRENE™ elastomers. Other suitable thermoplastic elastomer blends include butyl-30 polyethylene (e.g. in a ratio ranging between about 2:3 and about 3:2) and butyl-polypropylene.

Typically, the sealing ring and/or the second sealing ring additionally comprises lubricant material. Suitably, the sealing ring and/or the second sealing ring comprises35 up to 30%, preferably from 5 to 20% lubricant material.

In addition, the stem may also comprise lubricant material. Suitably, the valve stem comprises up to 30%, preferably from 5 to 20% lubricant material.

The term 'lubricant' herein means any material which reduces friction between the valve stem and seal. Suitable lubricants include silicone oil or a fluorocarbon polymer such as polytetrafluoroethane (PTFE) or fluoroethylene propylene (FEP).

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Lubricant can be applied to the stem, sealing ring or a second sealing ring by any suitable process including coating and impregnation, such as by injection or a tamponage process.

- 10 Optionally a moisture absorbing means may be incorporated within the dispenser as a component thereof. Alternatively, the moisture absorbing means may be a separate component of the formulation contained within the dispenser.

- 15 The moisture absorbing means may comprise a component or accessory for use with a canister or valve including a desiccant such as a molecular sieve, silica gel and/or a component or accessory that is made from a plastics material which is a natural desiccant, such as a polyamide, for example nylon, or may be moulded from other plastics material such as acetal or PBT. Alternatively, or in addition, the moisture absorbing means may comprise an internal lining or coating.
- 20 the moisture absorbing means may be incorporated into a treatment or coating for canisters and/or valves for preventing drug deposition and/or maintaining dose uniformity.

- 25 Other vapour or moisture absorbing materials include desiccants made from inorganic materials such as zeolites and aluminas. Such inorganic materials have high water absorption capacities and favourable water absorption isotherm shapes. The water absorption capacity of such materials typically varies from 20 to 50 weight percent.

- 30 Other exemplary moisture absorbing materials include, but are not limited to, alumina, bauxite, anhydrous calcium sulphate, water-absorbing clay, activated bentonite clay, molecular sieves, or other like materials.

- 35 In conjunction with the desiccant an additional compound may be added to act as a conduit/channelling agent to increase/optimize the efficiency of the moisture absorption properties. Such materials may include compounds such as polyethylene glycols.

Preferably, the means for absorbing moisture reduces the rise in moisture content over time, and/or the decrease in Fine Particulate Mass (FPM) over time by between 20 and 100%, for example, 40 to 70%, e.g. 45 to 55%. Typically, the component or accessory takes the form of a cap and/or a seal and/or a lining.

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Furthermore desiccant material may be included in the packaging material for the device.

10 The desiccant should be present in an amount sufficient to absorb any increases in moisture around the valve area of the MDI and thus alleviate or substantially prevent moisture increases inside the canister.

Typically, 100µg to 5g, for example, 1mg to 1g, e.g. 100mg to 500mg, such as about 100mg to 250mg of desiccant may be included.

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Canisters according to the invention may be filled with pharmaceutical formulations suitable for administration to patients e.g. those suffering from respiratory disorders such as asthma and COPD. Suitable formulations generally comprise a liquefied propellant gas such as a hydrofluoroalkane e.g. HFA134a or HFA227 and one or more drug substances.

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Thus provided is a canister according to the invention containing a formulation comprising a medicament and a hydrofluoroalkane propellant. Preferably the propellant is selected from 1,1,1,2-tetrafluoroethane, 1,1,1,2,3,3,3-heptafluoro-n-propane and mixtures thereof, especially 1,1,1,2-tetrafluoroethane.

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Preferably the drug will be in particulate form of a mass median diameter so as to permit inhalation into the bronchial airways i.e. generally less than 100 microns e.g. between around 1 and 10 microns especially 1-5 microns. Particle size reduction can be achieved e.g. by micronisation. Optionally the formulation may contain a dispersing agent and/or a co-solvent such as a C<sub>2-6</sub> aliphatic alcohol (e.g. ethanol) or a polyol (e.g. propylene glycol or polyethyleneglycol), especially ethanol. A suitable amount of dispersing agent would be around 0.001-50%, e.g. 0.05-5% by weight of drug. A suitable amount of co-solvent would be around 0.005-15% e.g. 0.1-5% by weight of formulation. When conventional dispersing agents such as lecithin, sorbitan trioleate and oleic acid are employed then ethanol will generally be employed as well. Alternatively, the drug may be dissolved in the liquefied propellant gas, generally by use of a co-solvent such as a C<sub>2-6</sub> aliphatic alcohol (e.g. ethanol) or a polyol (e.g.

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propylene glycol or polyethyleneglycol), especially ethanol. Certain preferred suspension formulations are free of all excipients besides the propellant and the particulate drug. Such preferred formulations consist only of particulate drug and propellant.

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Example medicaments may thus be selected from, for example, analgesics, e.g. codeine, dihydromorphine, ergotamine, fentanyl or morphine; anginal preparations, e.g. diltiazem; antiallergics, e.g. cromoglycate (e.g. as the sodium salt), ketotifen or nedocromil (e.g. as the sodium salt); anti-infectives e.g. cephalosporins, penicillins, streptomycin, sulphonamides, tetracyclines and pentamidine; antihistamines, e.g. methapyrilene; anti-inflammatories, e.g. beclomethasone (e.g. as the dipropionate ester), fluticasone (e.g. as the propionate ester), flunisolide, budesonide, rofleponide, mometasone e.g. as the furoate ester), ciclesonide, triamcinolone (e.g. as the acetone) or  $6\alpha$ ,  $9\alpha$ -difluoro- $11\beta$ -hydroxy- $16\alpha$ -methyl-3-oxo- $17\alpha$ -propionyloxy-androsta-1,4-diene- $17\beta$ -carbothioic acid S-(2-oxo-tetrahydro-furan-3-yl) ester; antitussives, e.g. noscapine; bronchodilators, e.g. albuterol (e.g. as free base or sulphate), salmeterol (e.g. as xinafoate), ephedrine, adrenaline, fenoterol (e.g. as hydrobromide), formoterol (e.g. as fumarate), isoprenaline, metaproterenol, phenylephrine, phenylpropanolamine, pirbuterol (e.g. as acetate), reproterol (e.g. as hydrochloride), rimeterol, terbutaline (e.g. as sulphate), isoetharine, tulobuterol or 4-hydroxy-7-[2-[[3-(2-phenylethoxy)propyl]sulfonyl]ethyl]amino]ethyl-2(3H)-benzothiazolone, diuretics, e.g. amiloride; anticholinergics, e.g. ipratropium (e.g. as bromide), tiotropium, atropine or oxitropium; hormones, e.g. cortisone, hydrocortisone or prednisolone; xanthines, e.g. aminophylline, choline theophyllinate, lysine theophyllinate or theophylline. It will be clear to a person skilled in the art that, where appropriate, the medicaments may be used in the form of salts, (e.g. as alkali metal or amine salts or as acid addition salts) or as esters (e.g. lower alkyl esters) or as solvates (e.g. hydrates) to optimise the activity and/or stability of the medicament and/or to minimise the solubility of the medicament in the propellant.

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Preferred medicaments are selected from albuterol, salmeterol, fluticasone propionate and beclomethasone dipropionate and salts or solvates thereof, e.g. the sulphate of albuterol and the xinafoate of salmeterol and mixtures thereof.

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Medicaments can also be delivered in combinations. Preferred formulations containing combinations of active ingredients contain salbutamol (e.g. as the free base or the sulphate salt) or salmeterol (e.g. as the xinafoate salt) or formoterol (e.g. as the fumarate salt) in combination with an anti-inflammatory steroid such as a

beclomethasone ester (e.g. the dipropionate) or a fluticasone ester (e.g. the propionate) or budesonide. A particularly preferred combination is a combination of fluticasone propionate and salmeterol, or a salt thereof (particularly the xinafoate salt). A further combination of particular interest is budesonide and formoterol (e.g. as the fumarate salt).

The final drug concentration in the formulation is preferably between 0.005-10% w/w more preferably 0.005-5% e.g. 0.01-1% w/w by weight of formulation. Preferably the concentration will be such as to deliver a therapeutic dose of the medicament in one or two actuations. Example therapeutic doses are 25, 50, 125 or 250µg per actuation of fluticasone propionate taken twice per day, 25µg per actuation of salmeterol (as xinafoate) taken twice per day, 100µg of salbutamol (e.g. as free base or sulphate) taken twice as needed and 50 or 250 µg per actuation of beclomethasone dipropionate taken twice two or three times per day.

The invention also provides use of canisters according to the invention in inhalation therapy, for example, for the treatment or prophylaxis of respiratory disorders; and use of a metered dose inhaler system in the treatment or prophylaxis of respiratory disorders are all alternative aspects of this invention.

A still further aspect of the present invention comprises a method of treating respiratory disorders such as, for example, asthma, which comprises administration by inhalation of an effective amount of a pharmaceutical formulation from a canister according to the invention.

The invention is illustrated by reference to the following examples.

#### Example 1

A laminate comprising two sheets of aluminium and a sheet of stainless steel each of thickness 0.2mm is formed using a pressure sensitive adhesive, heating and pressure lamination rollers, so as to form a sandwich of a steel layer between two aluminium layers. The laminate is deep drawn into the shape of a conventional MDI canister (approximate dimensions: diameter 22.1-22.2 mm, height 33.4-61.15mm; wall thickness 0.3-0.75 mm).

#### Example 2

The canister of Example 1 is spray coated on its internal surface with a blend of PTFE and PES and cured at between 350 to 400°C for approximately 10 minutes.

Example 3

The canister of Example 1 is spray coated on its internal surface with FEP and cured 350 to 400°C for approximately 10 minutes.

5 Example 4

The canisters of Examples 1, 2 and 3 may be fitted with a metering valve (Valois DF60) and filled through the valve with a suspension of fluticasone propionate in liquefied HFA134a.

10 Example 5

The canisters of Examples 1, 2 and 3 may be fitted with a metering valve (Valois DF60) and filled through the valve with a suspension of salmeterol xinafoate in liquefied HFA134a.

15 Example 6

The canisters of Examples 1, 2 and 3 may be fitted with a metering valve (Valois DF60) and filled through the valve with a suspension of salbutamol sulphate in liquefied HFA134a.

20 Example 7

The canisters of Examples 1, 2 and 3 may be fitted with a metering valve (Valois DF60) and filled through the valve with a suspension of fluticasone propionate and salmeterol xinafoate in liquefied HFA134a.

25 Throughout the specification and the claims which follow, unless the context requires otherwise, the word 'comprise', and variations such as 'comprises' and 'comprising', will be understood to imply the inclusion of a stated integer or step or group of integers but not to the exclusion of any other integer or step or group of integers or steps.

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## Claims

1. A canister suitable for use in metered dose inhalers and fitted with a metering valve characterised in that its walls are formed of a laminate comprising a first layer which is composed of a metal and a second layer which is composed of a strengthening material.
2. A canister as claimed in claim 1 wherein the second layer of strengthening material is a plastics material selected from: polyester, polypropylene, acetonitrile-styrene copolymer, high-impact styrene, nylon or polyacetal, optionally reinforced with a material selected from carbon fibres, filaments of glass, metal, boron, aluminium silicate, calcium carbonate, talc and barium sulphate, and wherein the first metal layer is composed of a material selected from aluminium and steel and characterised in that the first metal layer forms the internal surface of the canister and is optionally coated with one or more coating layers and an adhesive layer is optionally used between the plastics material layer and the first metal layer.
3. A canister as claimed in claim 1 wherein the second layer of strengthening material is a plastics material selected from: polyester, polypropylene, acetonitrile-styrene copolymer, high-impact styrene, nylon or polyacetal optionally reinforced with material selected from carbon fibres, filaments of glass, metal, boron, aluminium silicate, calcium carbonate, talc and barium sulphate, and wherein the second layer is optionally coated with one or more coating layers and the first metal layer is composed of a material selected from aluminium and steel and characterised in that the first metal layer forms the external surface of the canister and is optionally coated with one or more coating layers and an adhesive layer is optionally used between the plastics material layer and the first metal layer.
4. A canister according to claim 2 or claim 3 wherein the plastics material is reinforced.
5. A canister according to claim 4 wherein the plastics material is reinforced with a material selected from carbon fibres and metal.
6. A canister according to claim 1 characterised in that its walls are formed of a laminate wherein the laminate comprises a first metal layer of material forming the external surface of the canister selected from the list of metals consisting of: aluminium, brass, copper, chromium, iron, tin and steel, optionally coated with one or more coating layers, and a second layer of material selected from the list of strengthening materials consisting of aluminium, brass, copper, chromium, chromium oxide, iron, tin and steel, optionally coated with one or more coating layers.
7. A canister according to claim 1 characterised in that its walls are formed of a laminate wherein the laminate comprises a first metal layer of material forming the internal surface of the canister selected from the list of metals consisting

of: aluminium, brass, copper, chromium, iron, tin and steel, optionally coated with one or more coating layers and a second layer of material selected from the list of strengthening materials consisting of aluminium, brass, copper, chromium, chromium oxide, iron, tin and steel, optionally coated with one or more coating layers.

- 5 8. A canister according to claims 1 to 7 wherein the first metal layer is aluminium or stainless steel.
9. A canister according to claim 8 wherein the first metal layer is aluminium.
- 10 10. A canister according to any one of claims 1 to 9 wherein canister is formed from a laminate comprising a layer of steel sandwiched between two layers of aluminium and optionally the internal and/or external aluminium walls are provided with one or more coating layers.
11. A canister according to any one of claims 1 to 10 containing a formulation comprising medicament and hydrofluoroalkane propellant.
- 15 12. A canister according to claim 11 wherein the propellant is selected from 1,1,1,2-tetrafluoroethane, 1,1,1,2,3,3,3-heptafluoro-n-propane and mixtures thereof.
13. A canister according to claim 12 wherein the propellant is 1,1,1,2-tetrafluoroethane.
- 20 14. A canister according to any one of claims 9 to 13 wherein the formulation is a suspension formulation free of all excipients besides propellant and the particulate drug.
15. A canister according to any one of claims 11 to 14 wherein the medicament is selected from fluticasone propionate, beclomethasone dipropionate, salmeterol, albuterol and salts or solvates thereof and mixtures thereof.
- 25 16. A canister according to claim 15 wherein the medicament is fluticasone propionate in combination with salmeterol xinafoate.

# INTERNATIONAL SEARCH REPORT

Int. Application No

PCT/GB 02/01088

**A. CLASSIFICATION OF SUBJECT MATTER**  
IPC 7 B65D83/38 A61M15/00

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
IPC 7 B65D A61M

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

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☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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- \*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*O\* document referring to an oral disclosure, use, exhibition or other means
- \*P\* document published prior to the international filing date but later than the priority date claimed

- \*T\* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- \*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- \*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- \* & \* document member of the same patent family

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## INTERNATIONAL SEARCH REPORT

International Application No

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International Application No  
PCT/GB 02/01088

| Patent document<br>cited in search report | Publication<br>date | Patent family<br>member(s) | Publication<br>date |
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**DOCUMENT-IDENTIFIER:** WO 2072449 A1  
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IN METERED DOSE  
INHALERS  
**PUBN-DATE:** September 19, 2002

**INVENTOR-INFORMATION:**

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**INT-CL (IPC):** B65D083/38 , A61M015/00

**EUR-CL (EPC):** B65D083/14

## **ABSTRACT:**

CHG DATE=20021101 STATUS=O>A  
canister suitable for use in metered dose  
inhalers and fitted with a metering valve

characterised in that its walls are formed of a laminate comprising a first layer which is composed of a metal and a second layer which is composed of a strengthening material.